

As siRNAs derive from such long double-stranded RNAs, one could expect siRNA precursors to constitutively activate the interferon response... except that the whole pathway seems to be inactive in mouse oocytes [15]. Identifying the mouse tissues devoid of an interferon response could prove helpful in the search for endogenous siRNA function.

It can be expected, at least, that one particular feature of siRNAs will not make that task easy: endogenous siRNAs are expressed at very low levels — much lower than typical miRNAs. This may be linked to the activity of siRNAs: as guides for the endonucleolytic cleavage of their targets, they act catalytically, whereas most animal miRNAs act stoichiometrically (the target needs to be bound by the miRNA to be repressed, until it is degraded).

These observations suggest an impressive, almost scary, possibility. After the miRNA era (where so many functions have been ascribed to miRNAs, in so many physiological processes), we may be entering the 'siRNA era'. How many biological pathways will involve siRNAs? As siRNAs can act catalytically, minute amounts of these novel regulators could have tremendous effects;

undetected small RNAs may lie behind unexplained phenomena. Clearly, the exploration of small regulatory RNAs is not over.

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Short RNAs: How Big Is This Iceberg?

Recent studies have reported the identification of piwi-associated RNAs (piRNAs) in *Drosophila* somatic cells. Interestingly, these piRNAs derive from the 3' untranslated regions of a subset of transcribed protein-coding genes and, experimentation suggests, might control the expression of other protein-coding transcripts. Studies of additional organisms support the new pathway's presence across animals.

Isidore Rigoutsos

The RNA revolution is upon us. The last decade and a half has witnessed a flurry of research activity revolving around RNA, a molecule that for many years was thought to play only ancillary roles in the workings of a cell. Aided by technological and scientific advances, the repertoire of non-coding RNAs, i.e. RNAs that do not code for proteins, has been expanding steadily [1].

A particular class of short RNAs, microRNAs, has so far commanded the lion's share of researchers' attention.

Approximately 22 nucleotides long, microRNAs have been shown to post-transcriptionally regulate their targets in a sequence-dependent manner [2]. Originally believed to act through the 3' untranslated region (3'UTR) of the targeted mRNAs, microRNAs were recently shown to target the mRNA's amino-acid coding sequence (CDS) as well [3,4]. Recently, piwi-associated RNAs (piRNAs), another intriguing class of short RNAs with lengths between 23 and 30 nucleotides, burst onto the stage [5–8]. To date, piRNAs have been reported

in *Drosophila*, human, mouse, rat, *Xenopus*, zebrafish and the worm, and their known numbers are in the hundreds of thousands.

Drosophila piRNAs deriving from heterochromatic loci that are rich in nested, truncated, or damaged repeat elements, termed piRNA 'clusters', have been among the better studied. Such piRNAs participate in the 'ping-pong' cycle, an auto-amplification loop that is conserved in many metazoans and constitutes an adaptive immune response that maintains genomic integrity by suppressing the expression of transposons and other repetitive sequences [9,10].

In addition to piRNAs generated via the ping-pong cycle, other piRNAs that do not depend on auto-amplification have also been reported. The specifics of their generation had remained elusive for a while, but recent studies by the laboratories of Mikiko and

Given that *tj*'s mRNA apparently serves two purposes, namely protein synthesis and piRNA production, is the mRNA pre-processed in any way prior to entering the primary piRNA pathway? The answer appears to be negative. Indeed, Northern blot analysis by the Siomi team [13] using probes targeting, respectively, *tj*'s CDS and 3'UTR revealed a single band of the same length, suggesting no

separation of the primary mRNA into segments.

Further analysis of data from recent studies of *Drosophila* mutants [14,15], and of newly generated data [11] in conjunction with RNAi studies [13], allowed the two teams to establish that *zucchini* is involved in the 3'UTR-piRNA pathway. Also, study of larval ovaries of *tj* mutants by the Siomi team [13] showed TJ to be required for *piwi* activation in *Drosophila* gonadal somatic cells. Notably, there are several potential Maf binding sites near *piwi*'s putative transcription start site: for some of them, chromatin immunoprecipitation experiments indicated a direct association with TJ (Figure 1).

Once produced and loaded on Piwi, the *tj*-derived piRNAs have the ability to suppress targets. By searching for targets with (near) complementarity to *tj*'s 3'UTR piRNAs, the Siomi team [13] was able to identify *fasIII* as one such candidate target gene. Interestingly, all three putative sites of interaction with the sequenced piRNAs were in *fasIII* introns (Figure 1). Quantitative RT-PCR analysis showed that *fasIII* is upregulated in *piwi*- and *tj*-mutants, supporting the conjecture that *fasIII* is a piRNA target.

However, are the above results a *tj*-specific or, perhaps, a *Drosophila*-specific peculiarity? A wide-ranging investigation by Lai's team [11] provides the answer. Analysis of public data and additional primary data that they generated revealed strong 3'UTR piRNA production in mouse testis as well. These mouse piRNAs were proportionally highest in pre-pachytene testis (10 days postpartum) wherein Mili was found to be the predominant carrier of 3'UTR-piRNAs, in agreement with earlier observations [16]. Notably, despite the difference in the carrier molecule, the identities of the gene sources of the Mili- and Miwi-loaded 3'UTR-piRNAs were relatively similar. Further analysis also allowed Lai's team to reclassify previously published piRNA clusters as 3'UTR-derived ones. The team also analyzed piRNAs from *X. tropicalis* eggs that associated with the Y12 monoclonal antibody and identified numerous 3'UTR-piRNAs there as well, albeit they were not as abundant as in the mouse and in *Drosophila*. The findings of the Lai group [11] together with corroborating results by the Siomi team [13] (obtained

by studying *piwi* expression in mouse *tj*-mutant testis) suggest conservation of the identified pathway across animals.

Do all 3'UTRs give rise to piRNAs? Interestingly, by studying sequenced reads in conjunction with gene expression data in OSS, Lai's team showed that high-expression is not a sufficient condition for the generation of piRNAs. Instead, they found preference for mRNA substrates belonging to select gene ontology (GO) categories such as post-transcriptional gene regulation, regulation of structure and organ development, morphogenesis, cell differentiation, and others. A similar analysis of mouse testis data revealed analogous GO biases.

These above results provide evidence in support of a previously reported, computationally derived framework that linked intergenic/intronic regions with exons through short motifs termed pyknons [17]. Pyknons are organism-specific, variable-length, statistically significant nucleotide motifs that were arrived at using purely computational means. By definition, the pyknons of a given genome occur multiple times in intergenic and/or intronic regions, as well as at least once in an exon. Despite a pyknon's multiple occurrences in a genome, not all instances of the motif necessarily coincide with retroelements [17]. They were originally a genomic curiosity [18] but after piRNAs identified by deep sequencing became available [5–8], it was apparent that pyknons had computationally presaged the existence of piRNAs, both in terms of length and sequence. At the same time pyknons predicted putative regulatory connections between exonic and non-exonic sequences, similar to that reported by the Siomi group [13]. Notably, several of the GO categories that were found enriched in the original pyknon analysis [17] agree with those identified by Lai's team [11] as associated with 3'UTR-piRNAs. Subsequent analysis showed that human and mouse pyknons are over-represented in the introns of genes belonging to specific GO categories, and that the categories are the same in the two genomes despite the absence of conservation of the underlying intronic sequences [19]. Additionally, several examples of pyknons and published piRNAs that are *antisense* to introns have

been reported [19], underscoring the connection between pyknons and piRNAs and in accordance with the putative *fasIII* findings by Siomi's team [13].

The pyknon framework anticipates that each mRNA region (5'UTR, CDS, 3'UTR) is associated (as source, target, or perhaps both) to distinguishable-by-sequence collections of short RNAs with each region being linked to distinct GO categories: glimpses of this are evident in the results generated by Lai's team [11]. If we now combine this multitude of connections with a previous report on the differential expression of a human pyknon containing genomic region in a disease context [20], it becomes increasingly likely that these findings by the Siomi and Lai teams [11–13] represent further evidence that the community may have stumbled upon the tip of a regulatory iceberg of potentially staggering proportions.

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